

## **Native PAGE**

**AIM:** To perform Native PAGE for nucleic acids (DNA/RNA).

### **Materials Required:**

- 10X TAE
- 1X TAE
- 40% or 30% Polyacrylamide
- Autoclaved MilliQ
- APS
- TEMED
- EtBr

### **Procedure:**

- Set up the apparatus before making the gel. Make sure there is no leakage.
- Take a 25ml beaker and add 1ml of 10X TAE.
- Add 2ml of 40% Polyacrylamide (2.66ml if 30% PAA) and make the solution up to 10ml with autoclaved Milli-Q.
- Add 80ul APS and 10ul TEMED. Mix the solution quickly and pour it into the cassette immediately.
- Let it set for 20-30 mins.
- Set up the plates along with the gel in the box within the assigned space. Make sure you have placed it properly so that there is no leakage.
- Fill the box with 1X TAE till the mark shown in the box.
- Load your samples.
- Make the appropriate negative and positive connections, set up the time and volt, and run the gel.
- After the running is completed, remove the gel from the cassette and put it into a solution containing 50ml water and 2ml EtBr.
- Visualise and analyse the gel.